AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1 (Previously Presented): A compound represented by the following formula (I) or a physiologically acceptable salt thereof, or a hydrate thereof:

$$R^1$$
 R^2
 S
 W^1
 N
 R^4
 W^2-Q
(I)

wherein, R^1 and R^2 each independently represent hydrogen atom, a halogen atom, hydroxyl group, a group of OZ_{1-6} (the group of OZ_{1-6} represents an alkyl group having 1-6 carbon atoms or a fluoroalkyl group having 1-6 carbon atoms, which bonds via the oxygen atom), a group of $S(O)_nZ_{1-4}$ (Z_{1-4} represents an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms or an alkylene group derived therefrom), a group of $N(R^{12})(R^{13})$ (R^{12} and R^{13} each independently represent hydrogen atom, an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms), a group of Z_{1-8} which may be substituted (Z_{1-8} represents an alkyl group having 1-8 carbon atoms), a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group, or a 4- to 7-membered saturated or

partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_nZ_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$;

 W^1 represents a group selected from the group consisting of -CH=CH-, -N(R¹²)CO-, -CON(R¹²)-, -CH₂O- and -CH₂CH₂- (each of the aforementioned groups binds to the thiazole ring at the left end);

R³ represents hydrogen atom, a halogen atom, hydroxyl group or an amino group; R⁴ represents a group selected from the group consisting of hydrogen atom, a group of -OZ₀₋₄R⁵ (Z₀₋₄ represents an alkylene group having 1-4 carbon atoms, a fluorinesubstituted alkylene group having 1-4 carbon atoms or a single bond, and R⁵ represents a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_nZ_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4}))$, a group of $-S(O)_nZ_{0-4}R^5$, a group of $-N(R^6)(R^7)$ { R^6 and R^7 each independently represent hydrogen atom or Z₁₋₄, or they may bind to each other to form a saturated or unsaturated 5- to 7-membered ring (the ring may contain one or two hetero atoms as ring constituting atoms), and R⁶ and R⁷ may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a

group of OCON(R¹²)(R¹³), a group of CON(R¹²)(R¹³), a group of N(R¹²)CON(R¹²)(R¹³), a group of Z_{1-4} , a group of OZ_{1-4} , a group OZ_{1-4} , a group of OZ_{1-4} , group of OZ_{1-4} , group of OZ_{1-4} , group of OZ_{1-4} , a group of OZ_{1-4} , group of

or 6-membered unsaturated heterocyclic group which may be substituted; W^2 represents a single bond or $-C(R^8)=C(R^9)-(R^8)$ and R^9 each independently represent hydrogen atom, a halogen atom, a lower alkyl group, an alkoxy group, cyano group, carboxyl group, hydroxymethyl group, cyanomethyl group, vinyl group or a group of $N(R^{12})(R^{13})$, Q represents an acidic group, and W^2 and Q may bind together to form vinylidenethiazolidinedione in E- or Z-configuration or an equivalent heterocyclic ring; m and n each independently represent an integer of 0 to 2, and q represents an integer of 0 to 3.

Claim 2 (Previously Presented): A medicament composition for eliminating resistance of a microorganism with acquired drug resistance, which comprises a compound represented by formula (I) according to claim 1 or a physiologically acceptable salt thereof as an active ingredient.

Claim 3 (Previously Presented): A medicament composition for enhancing effect of an antimicrobial agent, which comprises a compound represented by formula (I) according to claim 1 or a physiologically acceptable salt thereof as an active ingredient.

Claim 4 (Currently Amended): A pharmaceutical composition for preventive and/or therapeutic treatment of a microbial infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump[,] which comprises a compound represented by formula (I) according to claim 1 or a physiologically acceptable salt thereof together with an antimicrobial agent.

Claim 5 (Previously Presented): A compound represented by the following formula (I) or a physiologically acceptable salt thereof, or hydrate thereof

$$\begin{array}{c|c}
R^1 \\
R^2 \\
S \\
W^1 \\
W^2 \\
R^3 \\
O \\
W^2 \\
W^3 \\
(I)
\end{array}$$

wherein, R^1 and R^2 each independently represent hydrogen atom, a halogen atom, hydroxyl group, a group of OZ_{1-6} (the group of OZ_{1-6} represents an alkyl group having 1-6 carbon atoms or a fluoroalkyl group having 1-6 carbon atoms, which bonds via the oxygen atom), a group of $S(O)_nZ_{1-4}$ (Z_{1-4} represents an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms or an alkylene group derived therefrom), a group of $N(R^{12})(R^{13})$ (R^{12} and R^{13} each independently represent hydrogen atom, an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms), a group of Z_{1-8} which may be substituted (Z_{1-8} represents an alkyl group having 1-8 carbon atoms or a fluoroalkyl group having 1-8 carbon atoms), a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group, or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group

consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_nZ_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$;

 W^1 represents a group selected from the group consisting of -CH=CH-, -N(R¹²)CO-, -CON(R¹²)-, -CH₂O- and -CH₂CH₂- (each of the aforementioned groups binds to the thiazole ring at the left end);

R³ represents hydrogen atom, a halogen atom, hydroxyl group or an amino group: R⁴ represents a group selected from the group consisting of hydrogen atom, a group of -OZ₀₋₄R⁵ (Z₀₋₄ represents an alkylene group having 1-4 carbon atoms, a fluorinesubstituted alkylene group having 1-4 carbon atoms or a single bond, and R⁵ represents a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_nZ_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$, a group of $-S(O)_n Z_{0-4} R^5$, a group of $-N(R^6)(R^7) \{R^6 \text{ and } R^7 \text{ each } 1\}$ independently represent hydrogen atom or Z₁₋₄, or they may bind to each other to form a saturated or unsaturated 5- to 7-membered ring (the ring may contain one or two hetero atoms as ring constituting atoms), and R⁶ and R⁷ may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of $OCON(R^{12})(R^{13})$, a group of $CON(R^{12})(R^{13})$, a group of $N(R^{12})CON(R^{12})(R^{13})$, a group of Z_{1-4} , a group of OZ_{1-4} , a group $S(O)_nZ_{1-4}$, group of CH_2OH , a group of

 $(CH_2)_mN(R^{12})(R^{13})$, carboxyl group, cyano group, a group of $CO-Z_{1-4}(R^{10})-N(R^{12})(R^{13})$ (R^{10} is a substituent corresponding to a side chain on an amino acid carbon or a group of $-Z_{1-4}-R^{11}$ (R^{11} represents a substituent which forms a quaternary salt) and a group of $CO-Z_{1-4}-N(R^{12})(R^{13})$ (CH_2)q }, a 5- or 6-membered aryl group which may be substituted and a 5-

or 6-membered unsaturated heterocyclic group which may be substituted; W^2 represents a single bond or $-C(R^8)=C(R^9)-(R^8$ and R^9 each independently represent hydrogen atom, a halogen atom, a lower alkyl group, an alkoxy group, cyano group, carboxyl group, hydroxymethyl group, cyanomethyl group, vinyl group or a group of $N(R^{12})(R^{13})$, Q represents an acidic group, and W^2 and Q may bind together to form vinylidenethiazolidinedione in E- or Z-configuration or an equivalent heterocyclic ring; m and n each independently represent an integer of 0 to 2, and q represents an integer of 0 to 3; R^{14} represents hydrogen atom, Z_{1-4} , $Z_{1-4}R^5$ or

Z₁₋₄OR⁵; and X represents C-H and Y represents C-H or nitrogen atom.

Claim 6 (Currently Amended): A medicament composition for preventive and/or therapeutic treatment of a microbial infection by microorganisms selected from

Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump which comprises a compound represented by the formula (I) according to claim 1 or a physiologically acceptable salt thereof as an active ingredient.

Claim 7 (Currently Amended): A method for therapeutic treatment of a microbial infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump comprising administering to a mammal in need thereof a therapeutically effective amount of the composition

according to claim 6.

Claim 8 (Previously Presented): The method according to claim 7, further comprising administering at least one antimicrobial agent.

Claim 9 (Previously Presented) The method according to claim 8, wherein the at least one antimicrobial agent is simultaneously administered with the composition.

Claim 10 (Previously Presented): The method according to claim 8, wherein the at least one antimicrobial agent is separately administered from the composition.

Claim 11 (Previously Presented): The method according to claim 8, wherein the at least one antimicrobial agent is successively administered with the composition.

Claim 12 (Previously Presented): The method according to claim 7 wherein the mammal is a human.

Claim 13 (Previously Presented): A method for preventive treatment of a microbial infection comprising administering to a mammal a preventively effective amount of the composition according to claim 6.

Claim 14 (Previously Presented): The method according to claim 13, further comprising administering at least one antimicrobial agent.

Claim 15 (Previously Presented): The method according to claim 14, wherein the at least one antimicrobial agent is simultaneously administered with the composition.

Claim 16 (Previously Presented): The method according to claim 14, wherein the at least one antimicrobial agent is separately administered from the composition.

Claim 17 (Previously Presented): The method according to claim 14, wherein the at least one antimicrobial agent is successively administered with the composition.

Claim 18 (Previously Presented): The method according to claim 13 wherein the mammal is a human.

Claim 19 (Canceled)

Claim 20 (Currently Amended): A medicament composition for preventive and/or therapeutic treatment of a microbial infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump which comprises a compound represented by the formula (I) according to claim 5 or a physiologically acceptable salt thereof as an active ingredient.

Claim 21 (Canceled)

Claim 22 (Currently Amended): A method for therapeutic treatment of a microbial infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump comprising administering to a mammal in need thereof a therapeutically effective amount of the composition according to claim 20.

Claim 23 (Canceled)

Claim 24 (Previously Presented): The method according to claim 22, further comprising administering at least one antimicrobial agent.

Claim 25. (Currently Amended): A method for therapeutic treatment of a microbial infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump comprising administering to a mammal in need thereof a therapeutically effective amount of a composition comprising a compound represented by formula (I) or a physiologically acceptable salt thereof as an active ingredient and at least one antimicrobial agent

$$\begin{array}{c|c}
R^1 \\
R^2 \\
S \\
W^1 \\
W^1 \\
X \\
N \\
N \\
N \\
W^2 - Q
\end{array}$$
(I)

wherein, R¹ and R² each independently represent hydrogen atom, a halogen atom, hydroxyl group, a group of OZ₁₋₆ (the group of OZ₁₋₆ represents an alkyl group having 1-6 carbon atoms or a fluoroalkyl group having 1-6 carbon atoms, which bonds via the oxygen atom), a group of $S(O)_{n}Z_{1-4}$ (Z_{1-4} represents an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms or an alkylene group derived therefrom), a group of $N(R^{12})(R^{13})$ (R^{12} and R^{13} each independently represent hydrogen atom, an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms), a group of Z_{1-8} which may be substituted (Z_{1-8} represents an alkyl group having 1-8 carbon atoms or a fluoroalkyl group having 1-8 carbon atoms), a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group, or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ₁₋₄, a group of S(O)_nZ₁₋₄, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of CONH₂, a group of CONH(Z_{1-4}) and a group of CON(Z_{1-4})(Z_{1-4}); W¹ represents a group selected from the group consisting of -CH=CH-, -N(R¹²)CO-. -CON(R¹²)-, -CH₂O- and -CH₂CH₂- (each of the aforementioned groups binds to the

R³ represents hydrogen atom, a halogen atom, hydroxyl group or an amino group;

thiazole ring at the left end);

R⁴ represents a group selected from the group consisting of hydrogen atom, a group of -OZ₀₋₄R⁵ (Z₀₋₄ represents an alkylene group having 1-4 carbon atoms, a fluorinesubstituted alkylene group having 1-4 carbon atoms or a single bond, and R⁵ represents a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_0Z_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$, a group of $-S(O)_nZ_{0-4}R^5$, a group of $-N(R^6)(R^7)$ { R^6 and R^7 each independently represent hydrogen atom or Z₁₋₄, or they may bind to each other to form. a saturated or unsaturated 5- to 7-membered ring (the ring may contain one or two hetero atoms as ring constituting atoms), and R⁶ and R⁷ may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OCON(R^{12})(R^{13}), a group of CON(R^{12})(R^{13}), a group of N(R^{12})CON(R^{12})(R^{13}), a group of Z_{1-4} , a group of OZ_{1-4} , a group $S(O)_nZ_{1-4}$, group of CH_2OH , a group of $(CH_2)_mN(R^{12})(R^{13})$, carboxyl group, cyano group, a group of $CO-Z_{1.4}(R^{10})-N(R^{12})(R^{13})$ (R¹⁰ is a substituent corresponding to a side chain on an amino acid carbon or a group of $-Z_{1-4}-R^{11}$ (R^{11} represents a substituent which forms a quaternary salt) and a group of $CO - Z_{1-4} - N(R^{12})(R^{13})$ (CH₂)q

 $(CH_2)q$ }, a 5- or 6-membered aryl group which may be substituted and a 5- or 6-membered unsaturated heterocyclic group which may be substituted; W^2 represents a single bond or $-C(R^8)=C(R^9)-(R^8)$ and R^9 each independently represent

hydrogen atom, a halogen atom, a lower alkyl group, an aikoxy group, cyano group, carboxyl group, hydroxymethyl group, cyanomethyl group, vinyl group or a group of $N(R^{12})(R^{13})$, Q represents an acidic group, and W^2 and Q may bind together to form vinylidenethiazolidinedione in E- or Z-configuration or an equivalent heterocyclic ring; m and n each independently represent an integer of 0 to 2, and q represents an integer of 0 to 3; R^{14} represents hydrogen atom, an alkyl group having 1, 3 or 4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms, $Z_{1-4}R^5$ or $Z_{1-4}OR^5$; and X and Y each independently represent C-H or nitrogen atom.

Claim 26 (Currently Amended): A method for preventive treatment of a microbial infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump comprising administering to a mammal a preventively effective amount of the composition according to claim 20.

Claim 27 (Currently Amended): A method for preventive treatment of a microbial-infection by microorganisms selected from Pseudomonas aeruginosa and bacteria having a genetically homologous drug efflux pump comprising administering to a mammal a preventively effective amount of a composition comprising a compound represented by the formula (I) or a physiologically acceptable salt thereof as an active ingredient and at least one antimicrobial agent[.]

wherein, R¹ and R² each independently represent hydrogen atom, a halogen atom,

hydroxyl group, a group of OZ_{1-6} (the group of OZ_{1-6} represents an alkyl group having 1-6 carbon atoms or a fluoroalkyl group having 1-6 carbon atoms, which bonds via the oxygen atom), a group of $S(O)_nZ_{1-4}$ (Z_{1-4} represents an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms or an alkylene group derived therefrom), a group of $N(R^{12})(R^{13})$ (R^{12} and R^{13} each independently represent hydrogen atom, an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms), a group of Z_{1-8} which may be substituted (Z_{1-8} represents an alkyl group having 1-8 carbon atoms or a fluoroalkyl group having 1-8 carbon atoms), a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group, or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group $^{c_{c_1}}$ consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_nZ_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$);

 W^1 represents a group selected from the group consisting of -CH=CH-, -N(R¹²)CO-, -CON(R¹²)-, -CH₂O- and -CH₂CH₂- (each of the aforementioned groups binds to the thiazole ring at the left end);

R³ represents hydrogen atom, a halogen atom, hydroxyl group or an amino group;
R⁴ represents a group selected from the group consisting of hydrogen atom, a group of
-OZ₀₋₄R⁵ (Z₀₋₄ represents an alkylene group having 1-4 carbon atoms, a fluorinesubstituted alkylene group having 1-4 carbon atoms or a single bond, and R⁵
represents a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group or a
4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl

group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_nZ_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$, a group of $-S(O)_0Z_{0-4}R^5$, a group of $-N(R^6)(R^7)$ { R^6 and R^7 each independently represent hydrogen atom or Z₁₋₄, or they may bind to each other to form a saturated or unsaturated 5- to 7-membered ring (the ring may contain one or two hetero atoms as ring constituting atoms), and R⁶ and R⁷ may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OCON(R^{12})(R^{13}), a group of CON(R^{12})(R^{13}), a group of N(R^{12})CON(R^{12})(R^{13}), a group of Z_{1-4} , a group of OZ_{1-4} , a group $S(O)_nZ_{1-4}$, group of CH_2OH , a group of $(CH_2)_mN(R^{12})(R^{13})$, carboxyl group, cyano group, a group of $CO-Z_{1.4}(R^{10})-N(R^{12})(R^{13})$ (R¹⁰ is a substituent corresponding to a side chain on an amino acid carbon or a group of $-Z_{1-4}-R^{11}$ (R^{11} represents a substituent which forms a quaternary salt) and a group of $CO - Z_{1-4} - N(R^{12})(R^{13})$ (CH₂)q

(CH₂)q }, a 5- or 6-membered aryl group which may be substituted and a 5- or 6-membered unsaturated heterocyclic group which may be substituted; W² represents a single bond or -C(R⁸)=C(R⁹)- (R⁸ and R⁹ each independently represent hydrogen atom, a halogen atom, a lower alkyl group, an alkoxy group, cyano group, carboxyl group, hydroxymethyl group, cyanomethyl group, vinyl group or a group of N(R¹²)(R¹³)), Q represents an acidic group, and W² and Q may bind together to form vinylidenethiazolidinedione in *E*- or *Z*-configuration or an equivalent heterocyclic ring; m and n each independently represent an integer of 0 to 2, and q represents an integer of 0

to 3; R^{14} represents hydrogen atom, an alkyl group having 1, 3 or 4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms, $Z_{1-4}R^5$ or $Z_{1-4}OR^5$; and X and Y each independently represent C-H or nitrogen atom.

Claim 28 (Currently Amended): A method for inhibiting drug resistance acquisition due to a drug efflux pump comprising administering to a mammal in need thereof an effective amount to inhibit drug resistance acquisition due to a drug efflux pump of the composition according to claim 6.

Claim 29 (Previously Presented): The method according to claim 28 wherein the mammal is a human.

Claim 30 (Currently Amended): A method for inhibiting drug resistance acquisition due to a drug efflux pump comprising administering to a mammal in need thereof an effective amount to inhibit drug resistance acquisition due to a drug efflux pump of the composition according to claim 20.

Claim 31 (Previously Presented): The method according to claim 30 wherein the mammal is a human.

Claim 32 (Currently Amended): A method for inhibiting drug resistance acquisition due to a drug efflux pump comprising administering to a mammal in need thereof an effective amount to inhibit drug resistance acquisition due to a drug efflux pump of a composition comprising a compound represented by formula (I) or a physiologically acceptable salt thereof as an active ingredient

$$\begin{array}{c|c}
R^1 \\
R^2 \\
S \\
W^1 \\
W^1 \\
X \\
N \\
N \\
N \\
W^2 - Q
\end{array}$$
(I)

wherein, R¹ and R² each independently represent hydrogen atom, a halogen atom, hydroxyl group, a group of OZ₁₋₆ (the group of OZ₁₋₆ represents an alkyl group having 1-6 carbon atoms or a fluoroalkyl group having 1-6 carbon atoms, which bonds via the oxygen atom), a group of $S(O)_n Z_{1-4}$ (Z_{1-4} represents an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms or an alkylene group derived therefrom), a group of N(R¹²)(R¹³) (R¹² and R¹³ each independently represent hydrogen atom, an alkyl group having 1-4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms), a group of Z_{1-8} which may be substituted (Z_{1-8} represents an alkyl group having 1-8 carbon atoms or a fluoroalkyl group having 1-8 carbon atoms), a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group, or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ₁₋₄, a group of S(O)_nZ₁₋₄, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$;

 W^1 represents a group selected from the group consisting of -CH=CH-, -N(R¹²)CO-, -CON(R¹²)-, -CH₂O- and -CH₂CH₂- (each of the aforementioned groups binds to the thiazole ring at the left end);

R³ represents hydrogen atom, a halogen atom, hydroxyl group or an amino group;

R⁴ represents a group selected from the group consisting of hydrogen atom, a group of -OZ_{0.4}R⁵ (Z_{0.4} represents an alkylene group having 1-4 carbon atoms, a fluorinesubstituted alkylene group having 1-4 carbon atoms or a single bond, and R⁵ represents a 5- to 7-membered cyclic alkyl group, an aryl group, a heteroaryl group or a 4- to 7-membered saturated or partially saturated heterocyclic group (the cyclic alkyl group, aryl group, heteroaryl group and heterocyclic group may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of OZ_{1-4} , a group of $S(O)_{0}Z_{1-4}$, a group of $N(R^{12})(R^{13})$, a group of Z_{1-4} , carboxyl group, a group of CO_2Z_{1-4} , group of $CONH_2$, a group of $CONH(Z_{1-4})$ and a group of $CON(Z_{1-4})(Z_{1-4})$, a group of $-S(O)_n Z_{0-4} R^5$, a group of $-N(R^6)(R^7)$ { R^6 and R^7 each independently represent hydrogen atom or Z₁₋₄, or they may bind to each other to form. a saturated or unsaturated 5- to 7-membered ring (the ring may contain one or two hetero atoms as ring constituting atoms), and R⁶ and R⁷ may have one to three substituents selected from the group consisting of a halogen atom, hydroxyl group, a group of $OCON(R^{12})(R^{13})$, a group of $CON(R^{12})(R^{13})$, a group of $N(R^{12})CON(R^{12})(R^{13})$, a group of Z_{1-4} , a group of OZ_{1-4} , a group $S(O)_nZ_{1-4}$, group of CH_2OH , a group of $(CH_2)_mN(R^{12})(R^{13})$, carboxyl group, cyano group, a group of $CO-Z_{1-4}(R^{10})-N(R^{12})(R^{13})$ (R¹⁰ is a substituent corresponding to a side chain on an amino acid carbon or a group of -Z₁₋₄-R¹¹ (R¹¹ represents a substituent which forms a quaternary salt) and a group of $CO - Z_{1-4} - N(R^{12})(R^{13})$

(CH₂)q }, a 5- or 6-membered aryl group which may be substituted and a 5- or 6-membered unsaturated heterocyclic group which may be substituted;

 W^2 represents a single bond or $-C(R^8)=C(R^9)-(R^8)$ and R^9 each independently

represent hydrogen atom, a halogen atom, a lower alkyl group, an alkoxy group, cyano group, carboxyl group, hydroxymethyl group, cyanomethyl group, vinyl group or a group of $N(R^{12})(R^{13})$), Q represents an acidic group, and W^2 and Q may bind together to form vinylidenethiazolidinedione in E- or Z-configuration or an equivalent heterocyclic ring; m and n each independently represent an integer of 0 to 2, and q represents an integer of 0 to 3; R^{14} represents hydrogen atom, an alkyl group having 1, 3 or 4 carbon atoms or a fluoroalkyl group having 1-4 carbon atoms, $Z_{1-4}R^5$ or $Z_{1-4}OR^5$; and X and Y each independently represent C-H or nitrogen atom.

33 (Previously Presented): The method according to claim 32 wherein the mammal is a human.